

[illegible]

Examiner Initials*	Cite No. <sup>1</sup>	Foreign Patent Document	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	Footnote
		Country Code <sup>2</sup> -Number <sup>3</sup> -Kind Code <sup>5</sup>				
/SY/	B1	EP 0 427 939	05-22-1991	American Cyanamid Co.		
/SY/	B2	EP 0 438 311	07-24-1991	Merck & Co. Inc.		
/SY/	B3	EP 0 823 416	08-07-1996	Ajinomoto Co., Inc.		
/SY/	B4	WO 91/07430	05-30-1991	The Upjohn Co.		

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PTO/SB/08b Substitute for Form 1449B/PTO			Application Number		10/565,366
<b>INFORMATION DISCLOSURE STATEMENT BY APPLICANT</b>  (Use as many sheets as necessary)			Filing Date		January 23, 2006
			First Named Inventor		HARTUNG, Rolf
			Art Unit		1764
			Examiner Name		To be assigned
Sheet	2	of	3	Attorney Docket	7601/84486

# **NON PATENT LITERATURE DOCUMENTS**

Examiner Initials *	Cite No. <sup>1</sup>	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T <sup>2</sup>
/SY/	C1	International Search Report for PCT/EP2004/006654 filed June 19, 2004.	
	C2	International Preliminary Report on Patentability for PCT/EP2004/006654 filed June 19, 2004.	
	C3	ALEXANDER, <i>et al.</i> , "A Diastereoselective Synthesis of (2S, 3R, 4S)-2-Amino-1-cyclohexyl-6-methylheptane-3,4-diol, The Abbott Aminodiol," <i>Tetrahedron Letters</i> 37:1961-1964 (1996).	
	C4	BIRCH, <i>et al.</i> , "The Structure and Some Reactions of the Iron Tricarbonyl Complex of Thebaine," <i>J. Chem. Soc. C</i> 531 (1968).	
	C5	BLÁHA, <i>et al.</i> , "Stereoisomeric Chiral 2,9-Diazabicyclo[4.4.0]Decane-3-,10-Diones as Models of Dipeptide Grouping: Synthesis, X-Ray, IR, NMR, and CD Studies," <i>Coll. Czech. Chem. Commun.</i> 49:712-742 (1984).	
	C6	CLINGMAN, <i>et al.</i> , "Effect of Amines on Hydrogenolysis of Alkylphenols," <i>J. Org. Chem.</i> 23:276-280 (February 1958).	
	C7	CORRINGER, <i>et al.</i> , "CCK-B Agonist of Antagonist Activities of Structurally Hindered and Peptidase-Resistant Boc-CCK <sub>4</sub> Derivatives," <i>J. Med. Chem.</i> 36:166-172 (1993).	
	C8	DEVANT, <i>et al.</i> , "Steroselektive Aldolreaktion Mit Chiralen Sekundären Acetamiden," <i>Chem. Ber.</i> 119:2191-2207 (1986).	
	C9	EISLER, <i>et al.</i> , "Amino Acids and Peptides. LXV. Analogues of Oxytocin," <i>Coll. Czech. Chem. Commun.</i> 31:4563-4580 (1966).	
	C10	FAUSTINI, <i>et al.</i> , "Stereospecificity in the Transformation of $\alpha$ -Aminoacids into Fluororacids," <i>Tetrahedron Letters</i> 22:4533-4536 (1981).	
	C11	HAYASHI, <i>et al.</i> , "Chiral ( $\beta$ -Aminoalkyl)Phosphines. Highly Efficient Phosphine Ligands for Catalytic Asymmetric Grignard Cross-Coupling," <i>J. Org. Chem.</i> 48:2195-2202 (1983).	
↓	C12	HARRIS, <i>et al.</i> , "Structure of Ristocetin A: Configurational Studies of the Peptide," <i>J. Am. Chem. Soc.</i> 104:363-365 (1982).	

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# **INFORMATION DISCLOSURE STATEMENT BY APPLICANT**

(Use as many sheets as necessary)

Sheet 3 of 3

Application Number 10/565,366  
Filing Date January 23, 2006  
First Named Inventor HARTUNG, Rolf  
Art Unit 1764  
Examiner Name To be assigned  
Attorney Docket 7601/84486

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/SY	C13	HOEKSTRA, <i>et al.</i> , "Large-Scale Synthesis of Anticoagulant Decapeptide MDL 28050," <i>Tetrahedron</i> 48: 307-318 (1992).	
	C14	ISHIDA, <i>et al.</i> , "Micropeptins 88-A to 88-F, Chymotrypsin Inhibitors from the Cyanobacterium <i>Microcystis aeruginosa</i> (NIES-88)," <i>Tetrahedron</i> 54: 5545-5556 (1998).	
	C15	MINNAARD, <i>et al.</i> , "Synthesis of Enantiomerically Pure Cyclohexylglycine," <i>Synthetic Communications</i> 29(24): 4327-4332 (1999).	
	C16	PLATA, <i>et al.</i> , "The Stereospecific Preparation of an Hydroxyethylene Isotere Precursor via a Novel Piperidine-2,5-Dione Template," <i>Tetrahedron Letters</i> 32(30): 3623-3626 (1991).	
	C17	SCHUDA, <i>et al.</i> , "A Short and Efficient Synthesis of (3S, 4S)-4-[( <i>tert</i> -Butyloxycarbonyl)amino]-5-cyclohexyl-3-hydroxypentanoic Acid Ethyl Ester," <i>J. Org. Chem.</i> 53: 873-875 (1988).	
	C18	TAMURA, <i>et al.</i> , "Guanylpiperidine Peptidomimetics: Potent and Selective bis-Cation Inhibitors of Factor Xa," <i>Bioorg. Med. Chem. Lett.</i> 10(8): 745-749 (April 2000).	
✓	C19	TAMURA, <i>et al.</i> , "A Synthesis of Optically Active $\alpha$ -Cyclohexylglycine," <i>Synthetic Communications</i> 8(5): 345-351 (1978).	

Examiner Signature

/Shawquia Young/

Date Considered

08/17/2008

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